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Typed Drawing

Word Mark	TAXOL
Goods and Services	IC 005. US 018. G & S: anti-cancer preparations. FIRST USE: 19911106. FIRST USE IN COMMERCE: 19911106
Mark Drawing Code	(1) TYPED DRAWING
Serial Number	74125254
Filing Date	December 20, 1990
Filed ITU	FILED AS ITU
Published for Opposition	August 20, 1991
Registration Number	1689497
Registration Date	May 26, 1992
Owner	(REGISTRANT) Bristol-Myers Squibb Company CORPORATION DELAWARE 345 Park Avenue New York NEW YORK 10017
Attorney of Record	NADINE FLYNN
Type of Mark	TRADEMARK
Register	PRINCIPAL
Affidavit Text	SECT 15. SECT 8 (6-YR). SECTION 8(10-YR) 20020201.
Renewal	1ST RENEWAL 20020201
Live/Dead Indicator	LIVE

PTO HOME	TRADEMARK	TESS HOME	NEW USER	STRUCTURED	FREE FORM	BROWSER DICT	TOP	HELP	PREV LIST
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NEWS	3	SEP 09	CA/CAPlus records now contain indexing from 1907 to the present
NEWS	4	Jul 15	Data from 1960-1976 added to RDISCLOSURE
NEWS	5	Jul 21	Identification of STN records implemented
NEWS	6	Jul 21	Polymer class term count added to REGISTRY
NEWS	7	Jul 22	INPADOC: Basic index (/BI) enhanced; Simultaneous Left and Right Truncation available
NEWS	8	AUG 05	New pricing for EUROPATFULL and PCTFULL effective August 1, 2003
NEWS	9	AUG 13	Field Availability (/FA) field enhanced in BEILSTEIN
NEWS	10	AUG 15	PATDPAFULL: one FREE connect hour, per account, in September 2003
NEWS	11	AUG 15	PCTGEN: one FREE connect hour, per account, in September 2003
NEWS	12	AUG 15	RDISCLOSURE: one FREE connect hour, per account, in September 2003
NEWS	13	AUG 15	TEMA: one FREE connect hour, per account, in September 2003
NEWS	14	AUG 18	Data available for download as a PDF in RDISCLOSURE
NEWS	15	AUG 18	Simultaneous left and right truncation added to PASCAL
NEWS	16	AUG 18	FROSTI and KOSMET enhanced with Simultaneous Left and Right Truncation
NEWS	17	AUG 18	Simultaneous left and right truncation added to ANABSTR
NEWS	18	SEP 22	DIPPR file reloaded
NEWS	19	SEP 25	INPADOC: Legal Status data to be reloaded
NEWS	20	SEP 29	DISSABS now available on STN
NEWS EXPRESS		OCTOBER 01	CURRENT WINDOWS VERSION IS V6.01a, CURRENT MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP), AND CURRENT DISCOVER FILE IS DATED 23 SEPTEMBER 2003
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SESSION

FULL ESTIMATED COST

0.21

0.21

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FILE COVERS 1907 - 10 Oct 2003 VOL 139 ISS 16

FILE LAST UPDATED: 9 Oct 2003 (20031009/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s hyaluronan and methotrexate

2669 HYALURONAN

34 HYALURONANS

2672 HYALURONAN

(HYALURONAN OR HYALURONANS)

12478 METHOTREXATE

19 METHOTREXATES

12480 METHOTREXATE

(METHOTREXATE OR METHOTREXATES)

L1 7 HYALURONAN AND METHOTREXATE

=> d L1 1-7 ibib abs hitrn

L1 ANSWER 1 OF 7 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2003:173470 CAPLUS

DOCUMENT NUMBER: 138:198677

TITLE: Use of **hyaluronan** as a protective agent in chemotherapy for improved therapeutic protocols

INVENTOR(S): Brown, Tracey Jean; Fox, Richard Mark

PATENT ASSIGNEE(S): Meditech Research Limited, Australia

SOURCE: PCT Int. Appl., 96 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003018062	A1	20030306	WO 2002-AU1160	20020827
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,				

LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
 PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,
 UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD,
 RU, TJ, TM
 RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG,
 CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
 PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR,
 NE, SN, TD, TG

PRIORITY APPLN. INFO.: AU 2001-7302 A 20010827

AU 2001-9504 A 20011213

AB The invention relates to the field of chemotherapy of diseases, e.g. cell proliferation disorders including cancer. In particular, the invention discloses the use of **hyaluronan** (HA) as a protective agent in the treatment of subjects. HA is administered in conjunction with a chemotherapeutic agent to facilitate the prolonged administration of a dose of the chemotherapeutic agent to be administered to a subject. Owing to the protective effects of the HA, the dose of chemotherapeutic agent may be substantially higher than a generally accepted ED, which would otherwise be expected to cause unacceptable side effects in the subject.

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L1 ANSWER 2 OF 7 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2002:71908 CAPLUS

DOCUMENT NUMBER: 136:112640

TITLE: **Hyaluronan** as a cytotoxic agent, drug pre-sensitizer and chemo-sensitizer in the treatment of disease

INVENTOR(S): Brown, Tracey; Fox, Richard

PATENT ASSIGNEE(S): Mediatech Research Limited, Australia

SOURCE: PCT Int. Appl., 70 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002005852	A1	20020124	WO 2001-AU849	20010713
W:		AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM		
RW:		GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG		
GB 2368525	A1	20020508	GB 2002-4331	20010713
EP 1301209	A1	20030416	EP 2001-951219	20010713
R:		AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR		
AU 760404	B2	20030515	AU 2001-72202	20010713
US 2003180382	A1	20030925	US 2003-88774	20030313

PRIORITY APPLN. INFO.: AU 2000-8795 A 20000714

WO 2001-AU849 W 20010713

AB The present invention relates to the enhancement of bioavailability of chemotherapeutic agents for the treatment of disease. In particular the present invention relates to a method of enhancing the bioavailability of a chemotherapeutic agent comprising the step of administering to a subject in need thereof a therapeutically effective amt. of **hyaluronan**. The present invention also relates to the treatment of a drug resistant disease whereby the drug resistance is overcome or alleviated with the use

of **hyaluronan** either alone or in combination with a chemotherapeutic agent. One disease that is frequently affected by both cellular resistance and bioavailability problems is cancer. The present invention also provides a method of treating cancer cells comprising the step of administering to a patient in thereof a therapeutically effective amt. of **hyaluronan**.

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L1 ANSWER 3 OF 7 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2001:798086 CAPLUS
DOCUMENT NUMBER: 135:348866
TITLE: RHAMM peptide conjugates for drug targeting
INVENTOR(S): Woloski, B. Michael R.; Williams, Ashley Martin;
Sereda, Terrance Jimmy; Wiebe, Deanna June
PATENT ASSIGNEE(S): Cangene Corporation, Can.
SOURCE: PCT Int. Appl., 121 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001080899	A2	20011101	WO 2001-CA533	20010420
WO 2001080899	A3	20020906		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
EP 1274461	A2	20030115	EP 2001-923439	20010420
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR			
PRIORITY APPLN. INFO.:			US 2000-198613P	P 20000420
			WO 2001-CA533	W 20010420

OTHER SOURCE(S): MARPAT 135:348866

AB The present invention provides protein conjugates having a glucose-aminoglycan-targeting domain conjugated directly or indirectly to a therapeutically useful protein via chem. or peptidyl linkage. A conjugate of the invention is disclosed in which a **hyaluronan**-binding protein is a receptor for hyaluronic acid-mediated mobility (RHAMM). The protein conjugates selectively target certain tissues and organs and are useful for treating or preventing various physiol. and pathol. conditions. Methods of their use and prepn. are described.

L1 ANSWER 4 OF 7 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2001:545502 CAPLUS
DOCUMENT NUMBER: 135:117219
TITLE: Hapten-coagulation agent-antineoplastic agent combinations for treating neoplasms
INVENTOR(S): Yu, Baofa
PATENT ASSIGNEE(S): USA
SOURCE: PCT Int. Appl., 83 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001052868	A1	20010726	WO 2001-US1737	20010118
WO 2001052868	C2	20030116		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

US 2002044919	A1	20020418	US 2001-765060	20010117
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PRIORITY APPLN. INFO.: US 2000-177024P P 20000119

AB Methods are provided for treating neoplasms, tumors and cancers, using one or more haptens and coagulation agents or treatments, alone or in combination with other anti-neoplastic agents or treatments. Also provided are combinations, and kits contg. the combinations for effecting the therapy.

REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L1 ANSWER 5 OF 7 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2000:493422 CAPLUS

DOCUMENT NUMBER: 133:109985

TITLE: A composition and method for the enhancement of the efficacy of drugs

INVENTOR(S): Brown, Tracey

PATENT ASSIGNEE(S): Meditech Research Limited, Australia

SOURCE: PCT Int. Appl., 126 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000041730	A1	20000720	WO 2000-AU4	20000106

W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

EP 1140198	A1	20011010	EP 2000-902481	20000106
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R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO

JP 2002534484	T2	20021015	JP 2000-593339	20000106
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NZ 512676	A	20030131	NZ 2000-512676	20000106
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ZA 2001005492	A	20021003	ZA 2001-5492	20010703
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PRIORITY APPLN. INFO.: AU 1999-8131 A 19990113

AU 1999-3938 A 19991109

WO 2000-AU4 W 20000106

AB The present invention relates to the enhancement of the efficacy of drugs, and more particularly, with overcoming the resistance of cells or organisms to drugs. In particular, the present invention provides a method for enhancing the effectiveness of a cytotoxic or antineoplastic agent, comprising the step of co-administering said agent with

hyaluronan, wherein co-administration with **hyaluronan** enhances the agent's cancer cell-killing potential. There was an increase in 5-FU uptake by tumors when 5-FU was injected with hyaluronic acid.

REFERENCE COUNT: 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L1 ANSWER 6 OF 7 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 1999:316588 CAPLUS

DOCUMENT NUMBER: 130:320837

TITLE: Oligosaccharides reactive with **hyaluronan** -binding protein, monoclonal antibodies recognizing **hyaluronan**-binding protein, and use in cancer therapy

INVENTOR(S): Toole, Bryan P.; Banerjee, Shib D.

PATENT ASSIGNEE(S): Trustees of Tufts College, USA

SOURCE: U.S., 22 pp., Cont.-in-part of U.S. Ser. No. 899,249, abandoned.

CODEN: USXXAM

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5902795	A	19990511	US 1994-306150	19940914

PRIORITY APPLN. INFO.: US 1992-899249 19920616

AB **Hyaluronan**-binding protein (HABP) is expressed on the cell surface during tumor cell and endothelial cell migration and during capillary-like tubule formation. Monoclonal antibodies and **hyaluronan** oligosaccharides are described which specifically recognize HABP and can be used to (1) inhibit tumor growth by preventing tumor vascularization, (2) inhibit tumor cell migration and (3) image tumors.

REFERENCE COUNT: 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L1 ANSWER 7 OF 7 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 1999:169161 CAPLUS

DOCUMENT NUMBER: 131:17430

TITLE: Production and elimination of **hyaluronan** in rheumatoid arthritis patients: estimation with a loading test

AUTHOR(S): Torsteinsdottir, Ingunn; Groth, Torgny; Lindqvist, Ulla

CORPORATE SOURCE: Department of Clinical Chemistry, University Hospital, Uppsala, S-751 85, Swed.

SOURCE: Seminars in Arthritis and Rheumatism (1999), 28(4), 268-279

CODEN: SAHRBF; ISSN: 0049-0172

PUBLISHER: W. B. Saunders Co.

DOCUMENT TYPE: Journal

LANGUAGE: English

AB To evaluate the benefit of detg. the maximal elimination rate (Vmax) and the endogenous prodn. of **hyaluronan** (HYA) in relation to the basal HYA concn. (c0) in rheumatoid arthritis (RA) patients; and to evaluate the compatibility of a new model for HYA kinetics, taking renal elimination into sep. account in the overall clearance of HYA from the blood. The calcns. of prodn. and elimination of HYA were based on the HYA loading test, which was performed in 21 patients with RA and 15 healthy controls. A blood sample was drawn before the loading test, followed by an i.v. injection of HYA as a single bolus dose of 7.5 mg. Blood samples were taken regularly during the next 60 min. A theor. model with computational anal. of the data collected was used for calcg. HYA prodn.

and elimination. Patients with RA had significantly higher c0 than healthy controls, although in 10 of 21 patients c0 was within the normal range. The RA patients also had higher Vmax than healthy controls, but the difference was not significant. The calcd. prodn. of HYA was increased in RA patients and correlated with c0. The new model for HYA kinetics, in which the renal elimination was taken sep. into account, proved to be more compatible than the previous model. The HYA loading test can help det. whether the increased serum level of HYA in RA patients is due to a high prodn. or reduced elimination of HYA or both.

REFERENCE COUNT: 40 THERE ARE 40 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> s hyaluronan and paclitaxel

2669 HYALURONAN

34 HYALURONANS

2672 HYALURONAN

(HYALURONAN OR HYALURONANS)

5434 PACLITAXEL

15 PACLITAXELS

5434 PACLITAXEL

(PACLITAXEL OR PACLITAXELS)

L2 4 HYALURONAN AND PACLITAXEL

=> d L2 1-4 ibib abs hitrn

L2 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2001:808942 CAPLUS

DOCUMENT NUMBER: 137:43795

TITLE: Identification of small molecule binding sites within proteins using phage display technology

AUTHOR(S): Rodi, D. J.; Agoston, G. E.; Manon, R.; Lapceovich, R.; Green, S. J.; Makowski, L.

CORPORATE SOURCE: Department of Discovery Research, Entremed, Inc., Rockville, MD, 20850, USA

SOURCE: Combinatorial Chemistry and High Throughput Screening (2001), 4(7), 553-572

CODEN: CCHSFU; ISSN: 1386-2073

PUBLISHER: Bentham Science Publishers

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Affinity selection of peptides displayed on phage particles was used as the basis for mapping mol. contacts between small mol. ligands and their protein targets. Anal. of the crystal structures of complexes between proteins and small mol. ligands revealed that virtually all ligands of mol. wt. 300 Da or greater have a continuous binding epitope of 5 residues or more. This observation led to the development of a technique for binding site identification which involves statistical anal. of an affinity-selected set of peptides obtained by screening of libraries of random, phage-displayed peptides against small mols. attached to solid surfaces. A random sample of the selected peptides is sequenced and used as input for a similarity scanning program which calcs. cumulative similarity scores along the length of the putative receptor. Regions of the protein sequence exhibiting the highest similarity with the selected peptides proved to have a high probability of being involved in ligand binding. This technique has been employed successfully to map the contact residues in multiple known targets of the anticancer drugs **paclitaxel** (Taxol), docetaxel (Taxotere) and 2-methoxyestradiol and the glycosaminoglycan **hyaluronan**, and to identify a novel **paclitaxel** receptor [1]. These data corroborate the observation that the binding properties of peptides displayed on the surface of phage particles can mimic the binding properties of peptides in naturally occurring proteins. It follows directly that structural context is relatively unimportant for detg. the binding properties of these

disordered peptides. This technique represents a novel, rapid, high
resoln. method for identifying potential ligand binding sites in the
absence of three-dimensional information and has the potential to greatly
enhance the speed of development of novel small mol. pharmaceuticals.

REFERENCE COUNT: 79 THERE ARE 79 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2001:545502 CAPLUS

DOCUMENT NUMBER: 135:117219

TITLE: Hapten-coagulation agent-antineoplastic agent
combinations for treating neoplasms

INVENTOR(S): Yu, Baofa

PATENT ASSIGNEE(S): USA

SOURCE: PCT Int. Appl., 83 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001052868	A1	20010726	WO 2001-US1737	20010118
WO 2001052868	C2	20030116		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,
SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU,
ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

US 2002044919	A1	20020418	US 2001-765060	20010117
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PRIORITY APPLN. INFO.: US 2000-177024P P 20000119

AB Methods are provided for treating neoplasms, tumors and cancers, using one
or more haptens and coagulation agents or treatments, alone or in
combination with other anti-neoplastic agents or treatments. Also
provided are combinations, and kits contg. the combinations for effecting
the therapy.

REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2000:493422 CAPLUS

DOCUMENT NUMBER: 133:109985

TITLE: A composition and method for the enhancement of the
efficacy of drugs

INVENTOR(S): Brown, Tracey

PATENT ASSIGNEE(S): Meditech Research Limited, Australia

SOURCE: PCT Int. Appl., 126 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000041730	A1	20000720	WO 2000-AU4	20000106

W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU,
CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL,
IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA,

MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI,
 SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM,
 AZ, BY, KG, KZ, MD, RU, TJ, TM
 RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
 DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
 CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
 EP 1140198 A1 20011010 EP 2000-902481 20000106
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
 IE, SI, LT, LV, FI, RO
 JP 2002534484 T2 20021015 JP 2000-593339 20000106
 NZ 512676 A 20030131 NZ 2000-512676 20000106
 ZA 2001005492 A 20021003 ZA 2001-5492 20010703
 PRIORITY APPLN. INFO.: AU 1999-8131 A 19990113
 AU 1999-3938 A 19991109
 WO 2000-AU4 W 20000106

AB The present invention relates to the enhancement of the efficacy of drugs,
 and more particularly, with overcoming the resistance of cells or
 organisms to drugs. In particular, the present invention provides a
 method for enhancing the effectiveness of a cytotoxic or antineoplastic
 agent, comprising the step of co-administering said agent with
hyaluronan, wherein co-administration with **hyaluronan**
 enhances the agent's cancer cell-killing potential. There was an increase
 in 5-FU uptake by tumors when 5-FU was injected with hyaluronic acid.

REFERENCE COUNT: 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 1999:64680 CAPLUS
 DOCUMENT NUMBER: 130:115045
 TITLE: **Paclitaxel** compositions containing
 hyaluronic acid of a molecular weight of less than
 750.000 Da
 INVENTOR(S): Asculai, Samuel S.; Moore, Adrian
 PATENT ASSIGNEE(S): Hyal Pharmaceutical Corporation, Can.
 SOURCE: PCT Int. Appl., 17 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9902151	A1	19990121	WO 1998-CA660	19980708
W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
CA 2208924	AA	19990109	CA 1997-2208924	19970709
AU 9882031	A1	19990208	AU 1998-82031	19980708
PRIORITY APPLN. INFO.: CA 1997-2208924 A 19970709 WO 1998-CA660 W 19980708				

AB **Hyaluronan** is used to deliver effective dosage amts. of
paclitaxel to a patient which medicine is present in a dosage amt.
 much less than the usual amt. presently being used when treating a patient
 with cancer. Taxol at 2.5 mg/kg and **hyaluronan** at 7.5 mg/kg
 decreased the wt. of tumors in mice from 470 to 391 g.
 REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> s L1 and toxicity
 292913 TOXICITY
 10626 TOXICITIES
 296196 TOXICITY
 (TOXICITY OR TOXICITIES)
 L3 2 L1 AND TOXICITY

=> d L3 1-2 ibib abs hitrn

L3 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN
 ACCESSION NUMBER: 2003:173470 CAPLUS
 DOCUMENT NUMBER: 138:198677
 TITLE: Use of **hyaluronan** as a protective agent in
 chemotherapy for improved therapeutic protocols
 INVENTOR(S): Brown, Tracey Jean; Fox, Richard Mark
 PATENT ASSIGNEE(S): Meditech Research Limited, Australia
 SOURCE: PCT Int. Appl., 96 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003018062	A1	20030306	WO 2002-AU1160	20020827
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				

PRIORITY APPLN. INFO.: AU 2001-7302 A 20010827
 AU 2001-9504 A 20011213

AB The invention relates to the field of chemotherapy of diseases, e.g. cell proliferation disorders including cancer. In particular, the invention discloses the use of **hyaluronan** (HA) as a protective agent in the treatment of subjects. HA is administered in conjunction with a chemotherapeutic agent to facilitate the prolonged administration of a dose of the chemotherapeutic agent to be administered to a subject. Owing to the protective effects of the HA, the dose of chemotherapeutic agent may be substantially higher than a generally accepted ED, which would otherwise be expected to cause unacceptable side effects in the subject.

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN
 ACCESSION NUMBER: 2002:71908 CAPLUS
 DOCUMENT NUMBER: 136:112640
 TITLE: **Hyaluronan** as a cytotoxic agent, drug
 pre-sensitizer and chemo-sensitizer in the treatment
 of disease
 INVENTOR(S): Brown, Tracey; Fox, Richard
 PATENT ASSIGNEE(S): Meditech Research Limited, Australia
 SOURCE: PCT Int. Appl., 70 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002005852	A1	20020124	WO 2001-AU849	20010713
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
GB 2368525	A1	20020508	GB 2002-4331	20010713
EP 1301209	A1	20030416	EP 2001-951219	20010713
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR			
AU 760404	B2	20030515	AU 2001-72202	20010713
US 2003180382	A1	20030925	US 2003-88774	20030313
PRIORITY APPLN. INFO.:			AU 2000-8795	A 20000714
			WO 2001-AU849	W 20010713

AB The present invention relates to the enhancement of bioavailability of chemotherapeutic agents for the treatment of disease. In particular the present invention relates to a method of enhancing the bioavailability of a chemotherapeutic agent comprising the step of administering to a subject in need thereof a therapeutically effective amt. of **hyaluronan**. The present invention also relates to the treatment of a drug resistant disease whereby the drug resistance is overcome or alleviated with the use of **hyaluronan** either alone or in combination with a chemotherapeutic agent. One disease that is frequently affected by both cellular resistance and bioavailability problems is cancer. The present invention also provides a method of treating cancer cells comprising the step of administering to a patient in thereof a therapeutically effective amt. of **hyaluronan**.

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> s L2 and toxicity
292913 TOXICITY
10626 TOXICITIES
296196 TOXICITY
(TOXICITY OR TOXICITIES)

L4 0 L2 AND TOXICITY

=> s hyaluronan and fluorouracil
2669 HYALURONAN
34 HYALURONANS
2672 HYALURONAN
(HYALURONAN OR HYALURONANS)
14497 FLUOROURACIL
268 FLUOROURACILS
14510 FLUOROURACIL
(FLUOROURACIL OR FLUOROURACILS)

L5 5 HYALURONAN AND FLUOROURACIL

=> d L5 1-5 ibib abs hitrn

L5 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2003 ACS on STN
ACCESSION NUMBER: 2003:173470 CAPLUS
DOCUMENT NUMBER: 138:198677
TITLE: Use of **hyaluronan** as a protective agent in

INVENTOR(S): chemotherapy for improved therapeutic protocols
 Brown, Tracey Jean; Fox, Richard Mark
 PATENT ASSIGNEE(S): Meditech Research Limited, Australia
 SOURCE: PCT Int. Appl., 96 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003018062	A1	20030306	WO 2002-AU1160	20020827
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				

PRIORITY APPLN. INFO.: AU 2001-7302 A 20010827
 AU 2001-9504 A 20011213

AB The invention relates to the field of chemotherapy of diseases, e.g. cell proliferation disorders including cancer. In particular, the invention discloses the use of **hyaluronan** (HA) as a protective agent in the treatment of subjects. HA is administered in conjunction with a chemotherapeutic agent to facilitate the prolonged administration of a dose of the chemotherapeutic agent to be administered to a subject. Owing to the protective effects of the HA, the dose of chemotherapeutic agent may be substantially higher than a generally accepted ED, which would otherwise be expected to cause unacceptable side effects in the subject.

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2003:9651 CAPLUS

DOCUMENT NUMBER: 139:138568

TITLE: Dry film made of hylan to prevent adhesion between two healing tissue surfaces

AUTHOR(S): Balazs, Endre A.; Larsen, Nancy E.; Leshchiner, Edward A.; Boney, John D.; Mitlitski, Vadim; Parent, Edward G.; Whetstone, Julie L.

CORPORATE SOURCE: Matrix Biology Institute, Ridgefield, NJ, 07657, USA
 SOURCE: Hyaluronan, [Proceedings of the International Cellucon Conference], 12th, Wrexham, United Kingdom, 2000 (2002), Meeting Date 2000, Volume 2, 7-12. Editor(s): Kennedy, John F.

Woodhead Publishing Ltd.: Cambridge, UK.

CODEN: 69DKVZ; ISBN: 1-85573-570-9

DOCUMENT TYPE: Conference

LANGUAGE: English

AB When the epithelial cell layer covering two adjacent tissues is removed accidentally or intentionally during surgical procedures, the underlying connective tissue will grow together during the wound- healing process. Similarly, when two connective tissue surfaces not covered by endothelium but sepd. by elastoviscous fluid contg. high mol. wt. **hyaluronan** are wounded by trauma or during surgical procedures, they can grow together during the healing process. Such adhesion between two tissue surfaces may interfere with function and the excessive new connective tissue formed (scar tissue) may exert pressure on adjacent nerves, causing chronic pain. This paper describes the use of new formulations of dry

films contg. only hylan. In animal models, this film prevented adhesion formation between two tissue surfaces denuded from their mesothelial or epithelial cell cover. The most important property of this film after it is hydrated by tissue fluids was that it still adhered to the tissue surface, ensuring its stay in place. Thus, it functions as a barrier material, sepg. the healing tissues. The films do not cause inflammation or foreign body reaction and they do not interfere with the healing of adjacent tissues. These films successfully prevented adhesions between tissue surfaces in liver and cecal abrasion models in rat and uterine horn abrasion models in rabbits. Such films can also be used as delivery vehicles for various drugs, influencing them by combining their phys. barrier effect with regulation effects on the healing process.

REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2003:7593 CAPLUS

DOCUMENT NUMBER: 139:127537

TITLE: Anti-cancer activity of **hyaluronan**

AUTHOR(S): Filion, Mario C.; Menard, Sonia; Filion, Benoit; Roy, Julie; Reader, Stephanie; Phillips, Nigel C.

CORPORATE SOURCE: Bioniche Therapeutics Research Centre, Montreal, QC, H4P 2R2, Can.

SOURCE: Hyaluronan, [Proceedings of the International Cellucon Conference], 12th, Wrexham, United Kingdom, 2000 (2002), Meeting Date 2000, Volume 1, 419-427. Editor(s): Kennedy, John F. Woodhead Publishing Ltd.: Cambridge, UK. CODEN: 69DKVZ; ISBN: 1-85573-570-9

DOCUMENT TYPE: Conference

LANGUAGE: English

AB Although **hyaluronan** (HA) has been shown to modulate cellular proliferation in numerous cell types little is known about its effect on cancer cells. We have evaluated the anti-proliferative activity of HA with a mol. mass of 5.0-7.5.times.105 Da towards a wide range of cancer cell types. We have found that HA at low concns. (< 80 .mu.g/mL) inhibits, in a dose-dependent manner, the cellular proliferation of prostate cancer cells (LNCaP, PC-3, DU-145), bladder cancer cells (HT-1376, RT-4, T24 and UMUC-3), breast cancer cells (MCF-7), melanoma cells (B16-F1) and fibrosarcoma cells (HT-1080). The presence of a no. of escape mechanisms assocd. with cancer progression such as p53/p21 mutations, Rb-mutations, p16 deletion, Fas resistance, absence of caspase-3 and overexpression of P-glycoprotein did not affect the ability of HA to inhibit cancer cell growth. The inhibition of cancer cell proliferation appeared to be independent of the level of expression of the HA receptor CD44. Furthermore, we found that HA potentiated the anti-proliferative activity of anti-cancer agents based on nucleic acids (mycobacterial cell wall complex and Mycobacterium phlei DNA) and of chemotherapeutic drugs (5-fluorouracil, cisplatin and tamoxifen). Our data indicates that HA having a mol. mass of 5.0-7.5.times.105 Da has considerable potential for development either as a chemotherapeutic agent or as an adjunct to anti-cancer agents.

REFERENCE COUNT: 18 THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2002:71908 CAPLUS

DOCUMENT NUMBER: 136:112640

TITLE: **Hyaluronan** as a cytotoxic agent, drug pre-sensitizer and chemo-sensitizer in the treatment of disease

INVENTOR(S): Brown, Tracey; Fox, Richard

PATENT ASSIGNEE(S): Meditech Research Limited, Australia

SOURCE: PCT Int. Appl., 70 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002005852	A1	20020124	WO 2001-AU849	20010713
W:			AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM	
RW:			GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG	
GB 2368525	A1	20020508	GB 2002-4331	20010713
EP 1301209	A1	20030416	EP 2001-951219	20010713
R:			AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR	
AU 760404	B2	20030515	AU 2001-72202	20010713
US 2003180382	A1	20030925	US 2003-88774	20030313
PRIORITY APPLN. INFO.:			AU 2000-8795	A 20000714
			WO 2001-AU849	W 20010713

AB The present invention relates to the enhancement of bioavailability of chemotherapeutic agents for the treatment of disease. In particular the present invention relates to a method of enhancing the bioavailability of a chemotherapeutic agent comprising the step of administering to a subject in need thereof a therapeutically effective amt. of **hyaluronan**. The present invention also relates to the treatment of a drug resistant disease whereby the drug resistance is overcome or alleviated with the use of **hyaluronan** either alone or in combination with a chemotherapeutic agent. One disease that is frequently affected by both cellular resistance and bioavailability problems is cancer. The present invention also provides a method of treating cancer cells comprising the step of administering to a patient in thereof a therapeutically effective amt. of **hyaluronan**.

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 1995:341470 CAPLUS

DOCUMENT NUMBER: 123:9822

TITLE: Synthesis and properties of hyaluronic acid conjugated nucleic acid analogs-1: synthesis of deacetylhyaluronan and introduction of nucleic acid bases

AUTHOR(S): Wada, Takehiko; Chirachanchai, Suwabun; Izawa, Naoto; Inaki, Yoshiaki; Takemoto, Kiichi

CORPORATE SOURCE: Faculty of Engineering, Osaka University, Suita, 565, Japan

SOURCE: Journal of Bioactive and Compatible Polymers (1994), 9(4), 429-47

CODEN: JBCPEV; ISSN: 0883-9115

DOCUMENT TYPE: Journal

LANGUAGE: English

AB The conjugation of nucleic acid base with **hyaluronan** was achieved by using the activated ester of pentachlorophenyl trichloroacetate. The conditions of de-N-acetylation of sodium hyaluronic acid were studied. In low concns. of NaOH, the degree of deacetylation was 26%, while in 7.4N NaOH, the degree of deacetylation was 76% and the viscosity was 1.12 dL/g. Thymine and 5-fluorouracil bases were quant. conjugated to deacetylhyaluronan in 65% and 51%, resp. The

interaction of thymine **hyaluronan** conjugate with the
complementary base of polyadenylate showed a small hypochromicity.

=> s L5 and toxicity

292913 TOXICITY

10626 TOXICITIES

296196 TOXICITY

(TOXICITY OR TOXICITIES)

L6 2 L5 AND TOXICITY

=> d L6 1-2 ibib abs hitrn

L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2003:173470 CAPLUS

DOCUMENT NUMBER: 138:198677

TITLE: Use of **hyaluronan** as a protective agent in
chemotherapy for improved therapeutic protocols

INVENTOR(S): Brown, Tracey Jean; Fox, Richard Mark

PATENT ASSIGNEE(S): Meditech Research Limited, Australia

SOURCE: PCT Int. Appl., 96 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003018062	A1	20030306	WO 2002-AU1160	20020827

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,
UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD,
RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG,
CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR,
NE, SN, TD, TG

PRIORITY APPLN. INFO.: AU 2001-7302 A 20010827

AU 2001-9504 A 20011213

AB The invention relates to the field of chemotherapy of diseases, e.g. cell
proliferation disorders including cancer. In particular, the invention
discloses the use of **hyaluronan** (HA) as a protective agent in
the treatment of subjects. HA is administered in conjunction with a
chemotherapeutic agent to facilitate the prolonged administration of a
dose of the chemotherapeutic agent to be administered to a subject. Owing
to the protective effects of the HA, the dose of chemotherapeutic agent
may be substantially higher than a generally accepted ED, which would
otherwise be expected to cause unacceptable side effects in the subject.

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2002:71908 CAPLUS

DOCUMENT NUMBER: 136:112640

TITLE: **Hyaluronan** as a cytotoxic agent, drug
pre-sensitizer and chemo-sensitizer in the treatment
of disease

INVENTOR(S): Brown, Tracey; Fox, Richard

PATENT ASSIGNEE(S): Meditech Research Limited, Australia

SOURCE: PCT Int. Appl., 70 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002005852	A1	20020124	WO 2001-AU849	20010713
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
GB 2368525	A1	20020508	GB 2002-4331	20010713
EP 1301209	A1	20030416	EP 2001-951219	20010713
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
AU 760404	B2	20030515	AU 2001-72202	20010713
US 2003180382	A1	20030925	US 2003-88774	20030313
PRIORITY APPLN. INFO.:		AU 2000-8795	A	20000714
		WO 2001-AU849	W	20010713

AB The present invention relates to the enhancement of bioavailability of chemotherapeutic agents for the treatment of disease. In particular the present invention relates to a method of enhancing the bioavailability of a chemotherapeutic agent comprising the step of administering to a subject in need thereof a therapeutically effective amt. of **hyaluronan**. The present invention also relates to the treatment of a drug resistant disease whereby the drug resistance is overcome or alleviated with the use of **hyaluronan** either alone or in combination with a chemotherapeutic agent. One disease that is frequently affected by both cellular resistance and bioavailability problems is cancer. The present invention also provides a method of treating cancer cells comprising the step of administering to a patient in thereof a therapeutically effective amt. of **hyaluronan**.

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